

Substitute for form 1449A PTO
(Revised 10/2001)

**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**
(Use as many sheets as necessary)

Sheet 1 of 2

Com. if Known

Application Number	10/762,078
Filing Date	January 21, 2004
First Named Inventor	Hardeman et al.
Group Art Unit	1623
Examiner Name	Not Assigned
Attorney Docket Number	047446/273072

U. S. PATENT DOCUMENTS

Examiner Initials*	Cite No.	Document Number Number - Kind Code (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages of Relevant Figures Appear
JEC	1	US-4,529,796	07-16-1985	Kang et al.	All
	2	US-4,988,680	01-29-1991	Halazy et al.	All
	3	US-5,536,822	07-16-1996	Haystead	All
	4	US-5,650,510	07-22-1997	Webb, II et al.	All
	5	US-5,780,617	07-14-1998	Van den Bosch et al.	All
	6	US-5,854,228	12-29-1998	Webb, II et al.	All

FOREIGN PATENT DOCUMENTS

Examiner Initials	Cite No.	Foreign Patent Document Country Code - Number Kind Code (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T
JEC	7	EP 0269947 B2	06-08-1988	Institute of Organic Chemistry and Biochemistry of the Academy of Sciences of the Czech Republic	All	
	8	EP 0338887 B1	10-25-1999	Merrell Dow Pharmaceuticals, Inc.	All	

OTHER DOCUMENTS

Examiner Initials	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T
JEC	9	BRESSI, J.C., et al., "Adenosine Analogues as Selective Inhibitors of Glyceraldehyde-3-phosphate Dehydrogenase of <i>Trypanosomatidae</i> via Structure-Based Drug Design," <i>J. Med. Chem.</i> 2001, pp. 2080-2093, Vol. 44	
	10	CHANG, Y.-T., et al., "Synthesis and Application of Functionally Diverse 2,6,9-Trisubstituted Purine Libraries as CDK Inhibitors," <i>Chemistry Biology</i> June 1999; pp. 361-375, Vol. 6, No. 6	

Examiner Signature	L. E. Crane	Date Considered	09/20/2006
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*Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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Sheet 2 of 2

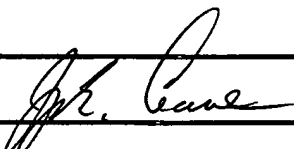
Complete if Known

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	11	HALBFINGER, E., et al., "Molecular Recognition of Modified Adenine Nucleotides by the P2Y ₁ -Receptor. 1. A Synthetic, Biochemical, and NMR Approach," <i>J. Med. Chem.</i> 1999, pp. 5325-5337, Vol. 42, No. 26	
	12	HERNANDEZ, A-I., et al., "Acyclic Nucleoside Analogues as Novel Inhibitors of Human Mitochondrial Thymidine Kinase," <i>J. Med. Chem.</i> 2002, pp. 4254-4263, Vol. 45, No. 19	
	13	HURYN, D.M., et al., "Synthesis of ISO-DDA, Member of a Novel Class of Anti-HIB Agents," <i>Tetrahedron Letters</i> 1989, pp. 6259-6262, Vol. 30, No. 46, Printed in Great Britain by Pergamon Press plc	
	14	VAN TILBURG, E.W., et al., "N ⁶ , 5'-Disubstituted Adenosine Derivatives as Partial Agonists for the Human Adenosine A ₃ Receptor," <i>J. Med. Chem.</i> 1999, pp. 1393-1400, Vol. 42, No. 8	

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Examiner Signature	I. E. Crane 	Date Considered	09/20/2006
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Sheet 1 of 1

Application Number 10/762,078
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First Named Inventor Hardeman et al.
Group Art Unit 1623
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Attorney Docket Number 047446/273072

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OTHER DOCUMENTS			
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	15	GUILLER, F., et al., "Linkers and Cleavage Strategies in Solid-Phase Organic Synthesis and Combinatorial Chemistry," <i>Chem. Rev.</i> 2000, pp. 2091-2157, Vol. 100, No. 6 (XP-002248255)	
	16	KUMAR, P., et al., "A Versatile Solid Phase Method for the Synthesis of Oligonucleotide-3'-Phosphates," <i>Tetrahedron Letters</i> , 1991, pp. 967-970, Vol. 32, No. 7 (XP-002112931)	
	17	Vyle, J.S., et al., "A Novel Solid Support for Synthesis of 2', 3'-Cyclic Phosphate Terminated Oligonucleotides," <i>Tetrahedron Letters</i> , 1998, pp. 7975, 7978, Vol. 39 (XP004137857)	

RTA01/2159960v1

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